PATENT 10/001,267 Docket 093/004p

CLAIM AMENDMENTS

+6504738654

1 to 12. CANCELLED

- 13. (Previously presented) A method for producing differentiated cells from primate pluripotent stem (pPS) cells, comprising:
 - a) obtaining a culture of pPS cells;
 - b) initiating differentiation of the pPS cells; and simultaneously or subsequently
 - c) culturing the cells of step b) in a medium containing a histone deacetylase inhibitor, butyrate until at least ~60% of the cultured cells have at least three of the following characteristics:
 - antibody-detectable expression of α₁-antitrypsin (AAT);
 - antibody-detectable expression of albumin;
 - absence of antibody-detectable expression of α-fetoprotein;
 - RT-PCR detectable expression of asialoglycoprotein receptor (ASGR);
 - evidence of glycogen storage;
 - evidence of cytochrome p450 activity;
 - · evidence of glucose-6-phosphatase activity; or
 - · the morphological features of hepatocytes.
- 14. (Previously presented) The method of claim 13, wherein at least about 60% of the cells have at least five of said characteristics.
- 15. (Previously presented) The method of claim 13, wherein at least about 80% of the cells have at least seven of said characteristics.

16 to 18. CANCELLED

19. (Previously presented) The method of claim 13, wherein differentiation of the pPS cells is initiated by forming embryoid bodies.

- 20. (Previously presented) The method of claim 13, wherein differentiation of the pPS cells is initiated by culturing in a medium containing dimethyl sulfoxide (DMSO), dimethylacetamide (DMA); hexmethylene bisacetamide, or another polymethylene bisacetamide.
- 21. (Previously presented) The method of claim 13, comprising further culturing the cells in a medium containing a cytokine or hormone selected from glucocorticoids, epidermal growth factor (EGF), insulin, TGF-α, TGF-β, fibroblast growth factor (FGF), hepatocyte growth factor (HGF), IL-1, IL-6, IGF-I, IGF-II, and HBGF-1.
- 22. (Previously presented) The method of claim 21, wherein the cells are cultured in a medium containing at least three of said cytokines or hormones.
- 23. (Previously presented) The method of claim 22, wherein the cells are cultured in a medium containing EGF, TGF-α, and HGF.
- 24. (Previously presented) The method of claim 13, further comprising maintaining the differentiated cells by culturing them in a medium containing a histone deacetylase inhibitor butyrate.
- 25. CANCELLED
- (Previously presented) The method of claim 27, wherein the pPS cells are human embryonic stem cells.
- 27. (Previously presented) A method for maintaining hepatocyte lineage cells in culture, comprising:
 - a) obtaining a population of cells differentiated from an established culture of primate pluripotent stem (pPS) cells, wherein at least ~60% of the differentiated cells have at least three of the following characteristics:
 - antibody-detectable expression of α₁-antitrypsin (AAT);
 - antibody-detectable expression of albumin;
 - absence of antibody-detectable expression of α-fetoprotein;
 - RT-PCR detectable expression of asialoglycoprotein receptor (ASGR);
 - evidence of glycogen storage;
 - evidence of cytochrome p450 activity;
 - evidence of glucose-6-phosphatase activity; or
 - the morphological features of hepatocytes; and then
 - b) culturing the differentiated cells in a medium containing a histone-deacetylase inhibitor, butvrate so that at least ~60% of the cultured cells maintain said characteristics.

PATENT 10/001,267 Docket 093/004p

- 28. (Previously presented) A method for producing differentiated cells from human embryonic stem (hES) cells, comprising:
 - a) obtaining a culture of hES cells;
 - b) initiating differentiation of the hES cells; and simultaneously or subsequently
 - c) culturing the cells of step b) in a medium containing a histone deacetylase inhibitor, butyrate until at least ~60% of the cultured cells have at least three of the following characteristics:
 - antibody-detectable expression of α₁-antitrypsin (AAT);
 - antibody-detectable expression of albumin;
 - absence of antibody-detectable expression of α-fetoprotein;
 - RT-PCR detectable expression of asialoglycoprotein receptor (ASGR);
 - evidence of glycogen storage;
 - evidence of cytochrome p450 activity;
 - · evidence of glucose-6-phosphatase activity; or .
 - the morphological features of hepatocytes.
- 29. (Previously presented) The method of claim 13, wherein the pPS cells are cultured with the histone deacetylase inhibitor butyrate without previously initiating differentiation.
- (Previously presented) The method of claim 13, wherein the pPS cells are cultured on an
 extracellular matrix without feeder cells before contact with the histone deacetylase inhibitor
 butyrate.
- 31. (Previously presented) The method of claim 28, wherein at least about 60% of the cells have at least five of said characteristics.
- 32. (Previously presented) The method of claim 28, wherein at least about 80% of the cells have at least seven of said characteristics.
- 33. CANCELLED
- 34. (Previously presented) The method of claim 28, comprising pre-differentiating the cells by culturing in a medium containing dimethyl sulfoxide (DMSO), dimethylacetamide (DMA); hexmethylene bisacetamide, or another polymethylene bisacetamide.
- 35. (Previously presented) The method of claim 28, comprising further culturing the cells in a medium containing at least three cytokines or hormones selected from glucocorticoids, epidermal

PATENT 10/001,267 Docket 093/004p

growth factor (EGF), insulin, TGF- α , TGF- β , fibroblast growth factor (FGF), hepatocyte growth factor (HGF), IL-1, IL-6, IGF-I, IGF-II, and HBGF-1.

- 36. (Previously presented) The method of claim 34, wherein the cells are cultured in a medium containing EGF, TGF-α, and HGF.
- 37. (Previously presented) The method of claim 27, wherein at least about 60% of the cells have at least five of said characteristics.
- 38. (*Previously presented*) The method of claim 27, wherein at least about 80% of the cells have at least seven of said characteristics.

39-40.	CANIC	T 1	Er
JJ-4U.	CAIVE	,ELL	

Upon allowance of the application, please renumber the claims as follows:

Claim	13	\rightarrow	1	Ciaim	28	\rightarrow	12
	14	\rightarrow	2		29	\rightarrow	4
	15	\rightarrow	3		30	\rightarrow	5
	19	\rightarrow	6		31	\rightarrow	13
	20	\rightarrow	7		32	\rightarrow	14
	21	 →	8		34	\rightarrow	15
	22	\rightarrow	9		35	→	16
	23	\rightarrow	10		36	\rightarrow	17
	24	\rightarrow	11		37	\rightarrow	19
	26	\rightarrow	21		38	\rightarrow	20
	27	\rightarrow	18				